

University of Groningen

ASYMMETRIC REDUCTIONS WITH A CHIRAL 1,4-DIHYDROPYRIDINE CROWN ETHER

DEVRIES, JG; KELLOGG, RM; Vries, J.G. de

Published in:
Journal of the American Chemical Society

DOI:
[10.1021/ja00504a060](https://doi.org/10.1021/ja00504a060)

IMPORTANT NOTE: You are advised to consult the publisher's version (publisher's PDF) if you wish to cite from it. Please check the document version below.

Document Version
Publisher's PDF, also known as Version of record

Publication date:
1979

[Link to publication in University of Groningen/UMCG research database](#)

Citation for published version (APA):
DEVRIES, JG., KELLOGG, RM., & Vries, J. G. D. (1979). ASYMMETRIC REDUCTIONS WITH A CHIRAL 1,4-DIHYDROPYRIDINE CROWN ETHER. *Journal of the American Chemical Society*, 101(10), 2759-2761. <https://doi.org/10.1021/ja00504a060>

Copyright

Other than for strictly personal use, it is not permitted to download or to forward/distribute the text or part of it without the consent of the author(s) and/or copyright holder(s), unless the work is under an open content license (like Creative Commons).

The publication may also be distributed here under the terms of Article 25fa of the Dutch Copyright Act, indicated by the "Taverne" license. More information can be found on the University of Groningen website: <https://www.rug.nl/library/open-access/self-archiving-pure/taverne-amendment>.

Take-down policy

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

Downloaded from the University of Groningen/UMCG research database (Pure): <http://www.rug.nl/research/portal>. For technical reasons the number of authors shown on this cover page is limited to 10 maximum.

Table I. Hyperfine Coupling Constants for PBN Adducts (in gauss)

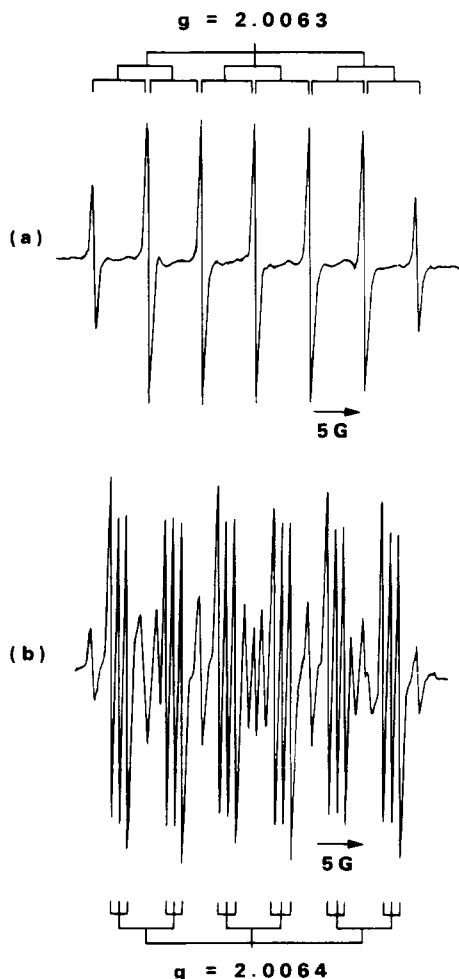
	A_N	A_H	A_D
H ₂ or D ₂ on ZnO	14.8	7.41	1.08
CO/H ₂ irradiation ^a	15.0	7.5	
irradiation of liquid alkanes ^b	14.8	7.0	

^a Reference 3. ^b Reference 2.

tained with the desorbed material permits the radical species to be identified (Figure 2). The spectrum obtained with adsorbed H₂ (trace a) is attributed to the addition of H• to I; the hyperfine pattern consists of a 1:1:1 triplet due to ¹⁴N, further split into 1:2:1 triplets due to two equivalent β protons. With D₂ (trace b), the addition product of D• to I gives a triplet-doublet-triplet splitting pattern, with the smaller triplet due to the β deuterium.

Figure 2 (b) also shows small amounts of the hydrogen addition product and another unidentified radical which was not present if a non-hydrogen-containing solvent such as carbon tetrachloride was used. A sample of ZnO was exchanged with D₂O and then outgassed in the usual way. Adsorption of hydrogen on this sample gave a spectrum showing the hydrogen addition product with a very small contribution from the deuterium addition product.

Table I compares the hyperfine splitting constants that we have observed with those reported in the literature for the addition products of H• and D• to I. The solution spectra are certainly those of II with R = H• or D•. The similarity of the adsorbed phase spectra to those from solution indicates that

**Figure 2.** EPR solution spectra of PBN adducts desorbed from ZnO containing (a) adsorbed H₂ and (b) adsorbed D₂.

the addition products are formed on the ZnO surface by reaction of I with adsorbed hydrogen or deuterium.

It has been shown by infrared spectroscopy that adsorption of H₂ on ZnO at room temperature involves a reversible dissociative chemisorption to form Zn-H and O-H species.^{5,6} Our experiments indicate that hydrogen adsorbed on ZnO can be abstracted by PBN. Further experiments are needed to determine which hydrogen is abstracted and to determine the mechanism of abstraction. Trapping of the adsorbed hydrogen by PBN does not necessarily imply the presence of free hydrogen atoms on the ZnO surface, but does indicate that the reactivity of adsorbed hydrogen resembles that of hydrogen atoms produced in radiolysis² or electrolysis⁴ experiments. The question of the extent of the radical character of hydrogen adsorbed on ZnO has still to be answered. Nevertheless, PBN is clearly a valuable spin trap to use in studying surface species having radical character. We envisage many systems of catalytic importance in which the presence of radical intermediates may be investigated by this technique.

References and Notes

- (1) E. G. Janzen, *Acc. Chem. Res.*, **4**, 31 (1971), and references therein.
- (2) S. A. Mao and L. Kevan, *J. Phys. Chem.*, **78**, 91 (1974).
- (3) S. Nagai, K. Matsuda, and M. Hatada, *J. Phys. Chem.*, **82**, 322 (1978).
- (4) P. H. Kasai and D. McLeod, *J. Phys. Chem.*, **82**, 619 (1978).
- (5) R. P. Eischens, W. P. Pliskin, and M. J. D. Low, *J. Catal.*, **1**, 180 (1962).
- (6) R. J. Kokes, A. L. Dent, C. C. Chang, and L. T. Dixon, *J. Am. Chem. Soc.*, **94**, 4429 (1972).

Taizo Uda, Akio Kazusaka
Russell F. Howe, George W. Kulks*

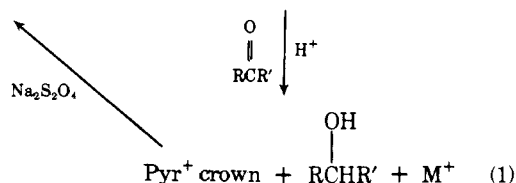
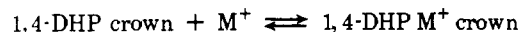
Laboratory for Surface Studies, Department of Chemistry
University of Wisconsin—Milwaukee
Milwaukee, Wisconsin 53201

Received November 6, 1978

Asymmetric Reductions with a Chiral 1,4-Dihydropyridine Crown Ether¹

Sir:

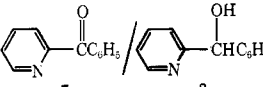
We are interested in the catalytic cycle shown in eq 1. A 1,4-dihydropyridine (DHP) is contained in a segment (for example a crown ether) capable of complexing a metal ion, M⁺. The encapsulated M⁺ then complexes with a carbonyl



compound, forming a ternary complex in which the carbonyl group is activated toward hydride acceptance through its complexation to M⁺.^{2,3} The pyridinium salt (Pyr⁺) formed on reduction of the carbonyl group is reduced back to 1,4-DHP with Na₂S₂O₄.⁴ Such a cycle has attractive synthetic and biomimetic aspects,⁵ especially if the 1,4-DHP-crown combination is chiral and is capable of carrying out reductions with a significant degree of asymmetric induction.⁶ We report here the preliminary results of work intended toward the achievement of the above goals.⁷

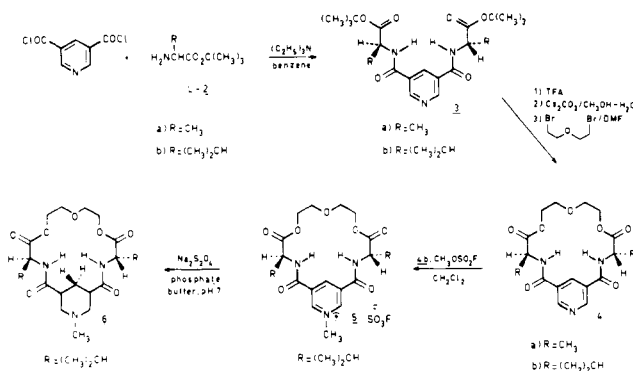
The synthetic route to the desired 1,4-DHP-crown compounds is shown in Scheme I. Chiral starting materials were the *tert*-butyl esters of optically pure L-alanine (**2a**) and L-

Table I

substrate/product	condi- tions (reaction time, h)	work- up proce- dure	NMR yield, % of alcohol (5, ClO ₄)	alcohol iso- lated yield, ^s %	[α] _D ^T , deg (concn)	[α] _D ^T , lit. deg (concn)	optical yield, %	absolute config of excess enantio- mer
 7 / 8 C ₆ H ₅ COCF ₃ (9) / C ₆ H ₅ - CHOHCF ₃ (10)	<i>a, b</i> (72)		65 (100)					
C ₆ H ₅ COCO ₂ C ₂ H ₅ (11) / C ₆ H ₅ - CHOHCO ₂ C ₂ H ₅ (12)	<i>b, c</i> (100)	<i>d</i>	82 (100)	58	[α] _D ²² + 9.7 (0.19) ^j	[α] _D ²⁴ + 14.2 <i>j.p.n</i>	68	<i>S</i>
C ₆ H ₅ COCONH ₂ (13) / C ₆ H ₅ CHOHCONH ₂ (14)	<i>c, e</i> (72)	<i>f, g</i>	80 (100)	61	[α] _D ²⁰ + 89.9 (0.76) ^k	[α] _D ²⁴ - 104 <i>k.m.o</i>	86	<i>S</i>
C ₆ H ₅ COCONH ₂ (13) / C ₆ H ₅ CHOHCONH ₂ (14)	<i>c, e</i> (48)	<i>f, h</i>	67 (100)	69	[α] _D ⁹ + 47.8 (0.55) ^l	[α] _D ⁹ + 74.7 <i>l.p.q</i>	64	<i>S</i>
C ₆ H ₅ COCONHC ₂ H ₅ (15) / C ₆ H ₅ - CHOHCONHC ₂ H ₅ (16)	<i>c, e</i> (75)	<i>f, i</i>	65 (100)	37	[α] _D ²⁰ + 26.5 (0.58) ^k	[α] _D ¹⁶ - 34.4 <i>k.m.r</i>	78	<i>S</i>

^aNMR experiment. ^b55 °C. ^c0.14 M in **6**, substrate, and Mg(ClO₄)₂·1.5H₂O in CH₃CN (CD₃CN). Reaction was carried out under N₂. ^dThe reaction mixture was concentrated, ether was added, and the precipitated **5** ClO₄ was filtered off. This procedure was repeated twice with the residue. Thereafter the combined ether extracts were concentrated, filtered, and subjected to preparative TLC (silica gel, C₆H₆). After Kugelrohr distillation the ¹H NMR spectrum of the alcohol **10** still showed some minor impurities. A sample was subjected to LC (silica gel, 82:18 CH₂Cl₂-hexane) and distilled again before measuring rotation. No impurities were detectable by either ¹H NMR spectroscopy or LC. ^eRoom temperature. ^fWater was added to the reaction mixture. After 30 min the solvent was evaporated; and the residue was dissolved in dry CH₃CN, filtered, and concentrated. ^gThe residue was subjected to preparative TLC (silica gel, 9:1 C₆H₆-(C₂H₅)₂O). Ethyl mandelate (**12**) was isolated and distilled in a Kugelrohr apparatus. No impurities in **12** could be detected either by ¹H NMR or LC. ^hThe alcohol was isolated by column chromatography (silica gel, 4:1 CH₂Cl₂-CH₃COCH₃) and further purified by LC (silica gel, 4:1 CH₂Cl₂-CH₃COCH₃). The compound was pure by LC and ¹H NMR and melted at 113.7–121.3 °C (lit. 122–122.5 °C for the pure enantiomer). ⁱThe alcohol was isolated by column chromatography (silica gel, 9:1 CH₂Cl₂-CH₃COCH₃) and further purified by LC (silica gel, 9:1 CH₂Cl₂-CH₃COCH₃), colorless oil (lit. mp 65.5–66.5 °C for the pure enantiomer), and was pure by both ¹H NMR and LC. ^jIn C₆H₆. ^kIn C₂H₅OH. ^lIn CH₃COCH₃. ^mFor optically pure material (*R*). ⁿReference 15. ^oReference 16. ^pFor optically pure material (*S*). ^qReference 17. ^rReference 18. ^sCorrected for recovered substrate.

Scheme 1



valine (**2b**). Reactions of these amino acid esters with **1** gave, respectively, **3a** (80% yield, mp 137.3–138.6 °C, [α]_D²¹ +26.8° (*c* 1.00, C₂H₅O₂CCH₃)) and **3b** (88% yield, mp 170.1–170.4 °C, [α]_D²¹ +37.7° (*c* 1.00, C₂H₅O₂CCH₃)). For the crucial ring-closure reaction **3a** and **3b** were first deblocked with trifluoroacetic acid (TFA) and were then converted to their dicesium salts and allowed to react with 1,5-dibromo-3-oxopentane in DMF. The cesium salt method, first developed for peptide chemistry, and known not to involve racemization,⁸ has recently been applied by us for the synthesis of crown ethers⁹ and macrocyclics.¹⁰ There was obtained **4a** (22% yield overall, 26% for ring closure, mp 259.1–262.3 °C dec, [α]_D²³ -94.0° (*c* 0.95, DMF)) and **4b** (32% yield overall, 48% for ring closure, mp 251.4–254.3 °C, [α]_D²⁰ -126.8° (*c* 1.02, DMF)). Thus far only **4b** has been used for further experiments. Methylation with methyl fluorosulfonate gave pyridinium salt **5** (80% yield, nonreproducible decomposition point, [α]_D²² -159.6° (*c* 1.00, CH₃OH)). Reduction at room temperature with sodium dithionite in a pH 7 phosphate buffer afforded **6** (98% yield, mp 140.5–142.4 °C, UV (CH₂Cl₂) 352 nm (ε 2500), [α]_D²⁵ -133.9° (*c* 1.04, CH₃CN)). This reduction method is a large improvement over literature procedures, which call for basic conditions;¹¹ excellent yields of very pure product are obtained in 10–20 min. Analytical data for all new compounds are good and spectral data are in accord with the

proposed structures. We believe that all pyridine compounds reported here are optically pure (note that inversion at one asymmetric center produces a meso diastereomer). As far as we are aware this is the first report of the synthesis of amino acid containing "crown ethers".

The results of several reductions with **6** are given in Table I. Reactions were carried out under nitrogen in acetonitrile with equimolar amounts of **6**, substrate, and Mg(ClO₄)₂·1.5H₂O.¹² In the presence of oxygen, **6** was completely oxidized to the perchlorate salt of **5** without detectable reduction of substrate. We observed a direct correlation between the sensitivity of **6** to oxidation and the ease of reduction of the substrate. In the absence of Mg(ClO₄)₂·1.5H₂O either with or without oxygen no significant oxidation of **6** or reduction of substrate occurred.

Reduction by **6** of 2-benzoylpyridine (**7**) to alcohol **8** proceeded only slowly (72 h) even at the relatively high temperature of 55 °C. Data for the maximum rotation of **8** as well as its absolute configuration have not been reported and therefore this reaction was not examined further. Optical data are, however, available for the alcohol **10** derived from trifluoroacetophenone (**9**), which is a popular substrate for reductions by 1,4-dihydropyridines. We found the reaction to be very sluggish, taking 5 days at 55 °C to go to completion. The degree of asymmetric induction in **10**, 68%, is, however, quite satisfying. The reductions of the ethyl ester (**11**) and amide (**13**) of phenylglyoxylate proceeded exceptionally smoothly. The optical inductions in **12** and **14** of, respectively, 86 and 64% are also very good indeed considering that the chiral centers in **6** are five bonds removed from the site of hydride donation.¹³ The reduction of the *N*-ethylamide **15** to alcohol **16** has been attended thus far by experimental difficulties: the isolated chemical yield is poor but the optical induction is high with the same stereochemical result (*S* configuration; see below) as observed in other reductions.

In all cases care was taken to avoid optical fractionation of the alcohol during workup.¹⁴ The recovered perchlorate of pyridinium salt **5** could be reduced by dithionite back to **6** with undiminished rotation. This completes, albeit in two separate reactions, the catalytic cycle of eq 1.

These results represent some of the highest asymmetric in-

ductions ever achieved with optically active 1,4-dihydropyridines.⁶ The consistent formation of an excess of the *S* enantiomer (the relative priorities of the groups are the same for all the optically active alcohols allowing direct comparison) strongly suggests structurally related transition states for reduction. ¹H NMR shielding effects in the presence of Mg²⁺ indicate complexation of Mg²⁺ close to the diethylene glycol bridge of **6**. Assuming that the oxygen of carbonyl group complexes to Mg²⁺ with the carbonyl carbon oriented toward the 4 position of the 1,4-dihydropyridine, that the phenyl substituent is the largest group, and that complexed α -dicarbonyl compounds assume a *cis* conformation for the carbonyl groups, the observed *S* configurations can be predicted. It is important to note that **6** is rather rigid owing to the two amide linkages.

Further experiments are in progress.¹⁹

References and Notes

- (1) Dedicated to Professor E. Havinga, University of Leiden, on the occasion of his 70th birthday.
- (2) See for examples of this and other recent approaches (a) S. Shinkai and T. C. Bruice, *Biochemistry*, **12**, 1750 (1973); (b) U. K. Pandit and F. R. MasCabr , *J. Chem. Soc. D*, 552 (1971); (c) U. K. Pandit, F. R. MasCabr , R. A. Gase, and M. J. de Nie-Sarink, *J. Chem. Soc., Chem. Commun.*, 627 (1974); (d) U. K. Pandit, R. A. Gase, F. R. MasCabr , and M. J. de Nie-Sarink, *ibid.*, 211 (1975); (e) R. A. Gase, G. Boxhoorn, and U. K. Pandit, *Tetrahedron Lett.*, 2889 (1976); (f) D. C. Dittmer, A. Lombardo, F. H. Batzold, and C. S. Greene, *J. Org. Chem.*, **41**, 2976 (1976); (g) Y. Ohnishi, M. Kagami, and A. Ohno, *J. Am. Chem. Soc.*, **97**, 4766 (1975); (h) Y. Ohnishi, M. Kagami, and A. Ohno, *Tetrahedron Lett.*, 2437 (1975); (i) Y. Ohnishi, T. Numakunai, T. Kimura, and A. Ohno, *ibid.*, 2699 (1976); (j) D. J. Creighton and D. S. Sigman, *J. Am. Chem. Soc.*, **93**, 6314 (1971); (k) R. A. Gase, G. Boxhoorn, and U. K. Pandit, *Tetrahedron Lett.*, 2889 (1976); (l) U. K. Pandit, P. C. Kelijzer, and R. A. Gase, *J. Chem. Soc., Chem. Commun.*, 493 (1976); (m) U. K. Pandit, H. van Dam, and J. B. Stevens, *Tetrahedron Lett.*, 913 (1977); (n) A. Ohno, T. Kimura, H. Yamamoto, S. G. Kim, S. Oka, and Y. Ohnishi, *Bull. Chem. Soc. Jpn.*, **50**, 1535 (1977); (o) A. Ohno, M. Ikeguchi, T. Kimura, and S. Oka, *J. Chem. Soc., Chem. Commun.*, 328 (1978); (p) Y. Ohnishi, M. Kagami, T. Numakunai, and A. Ohno, *Chem. Lett.*, 915 (1976); (q) T. Endo, Y. Hayashi, and M. Okawara, *ibid.*, 391 (1977).
- (3) For applications of NADH itself for synthetic and mechanistic purposes, see J. B. Jones and J. F. Beck in "Applications of Biochemical Systems in Organic Chemistry", Part I, J. B. Jones, C. J. Sih, and D. Perlman, Eds., "Techniques of Chemistry", Vol. X, Wiley-Interscience, New York, 1976, p 107.
- (4) (a) O. Warburg, W. Christian, and A. Guese, *Biochem. Z.*, **282**, 157 (1935); (b) K. Wallenfels and H. Schuly, *Justus Liebigs Ann. Chem.*, **621**, 106 (1959); (c) B. M. Anderson, C. J. Ciatti, and N. O. Kaplan, *J. Biol. Chem.*, **234**, 1219 (1959); (d) D. Mauzerall and F. H. Westheimer, *J. Am. Chem. Soc.*, **77**, 2261 (1955).
- (5) A reduction of thiones by such a catalytic cycle has been reported recently; K. Nakamura, A. Ohno, S. Yasui, and S. Oka, *Tetrahedron Lett.*, 4815 (1978).
- (6) For examples of asymmetric induction with 1,4-dihydropyridines, see (a) F. R. MasCabr , Dissertation, Amsterdam 1974; (b) ref 2g; (c) Y. Ohnishi, T. Numakunai, and A. Ohno, *Tetrahedron Lett.*, 3813 (1975); (d) ref 2i; (e) A. Ohno, H. Yamamoto, T. Kimura, and S. Oka, *Tetrahedron Lett.*, 4585 (1976); (f) K. Nishiyama, N. Baba, J. Oda, and Y. Inouye, *J. Chem. Soc., Chem. Commun.*, 101 (1976); (g) ref 2p; (h) A. Ohno, T. Kimura, S. G. Kim, H. Yamamoto, S. Oka, and Y. Ohnishi, *Bioorg. Chem.*, **6**, 21 (1977); (i) T. Makino, N. Baba, J. Oda, and Y. Inouye, *Chem. Ind. (London)*, 277 (1977); (j) H. J. van Ramesdonk, J. W. Verhoeven, U. K. Pandit, and Th. J. de Boer, *Recl. Trav. Chim. Pays-Bas*, **97**, 195 (1978); (k) see also ref 2o and 2q.
- (7) For examples of catalytically active crown ethers, see (a) Y. Chao and D. J. Cram, *J. Am. Chem. Soc.*, **98**, 1015 (1976); (b) T. Matsui and K. Koga, *Tetrahedron Lett.*, 1115 (1978); (c) T. J. van Bergen and R. M. Kellogg, *J. Am. Chem. Soc.*, **99**, 3882 (1977); (d) J. P. Behr and J.-M. Lehn, *J. Chem. Soc., Chem. Commun.*, 143 (1978); (e) J.-M. Lehn and C. Sirlin, *ibid.*, 949 (1978).
- (8) S.-S. Wang, B. F. Gisin, D. P. Winter, R. Makofske, J. D. Kulesha, C. Tzougraki, and J. Meienhofer, *J. Org. Chem.*, **42**, 1286 (1977), and references cited therein.
- (9) (a) O. Piepers and R. M. Kellogg, *J. Chem. Soc., Chem. Commun.*, 383 (1978); (b) B. J. van Keulen, R. M. Kellogg, and O. Piepers, *ibid.*, in press.
- (10) W. H. Kruizinga and R. M. Kellogg, *J. Chem. Soc., Chem. Commun.*, in press.
- (11) See, for example, J. G. de Vries, T. J. van Bergen, and R. M. Kellogg, *Synthesis*, 246 (1977).
- (12) G. F. Smith and E. G. Koch, *Z. Anorg. Chem.*, **223**, 17 (1935).
- (13) An optical induction of 100% has been reported for a case in which the hydride is donated from the optically active center.²⁰ The term "hydride donation" as used here is a formalism having no mechanistic implications.
- (14) Ohno²⁹ has demonstrated that no optical fractionation occurs during workup of alcohol **12**.
- (15) H. M. Peters, D. M. Feigl, and H. S. Mosher, *J. Org. Chem.*, **33**, 4245 (1968).
- (16) R. Roger, *J. Chem. Soc.*, 2168 (1932).
- (17) (a) H. Wren, *J. Chem. Soc.*, **95**, 1583 (1909); (b) "Dictionary of Organic Compounds", Vol. 4, Eyre and Spottiswoode, London, 1965, p 2051.
- (18) A. Mackenzie, G. Martin, and H. G. Rule, *J. Chem. Soc.*, **105**, 1583 (1914); ref 17b.
- (19) Previous publication in this series: R. H. van der Veen, R. M. Kellogg, A. Vos, and T. J. van Bergen, *J. Chem. Soc., Chem. Commun.*, 923 (1978).

J. G. de Vries, Richard M. Kellogg*

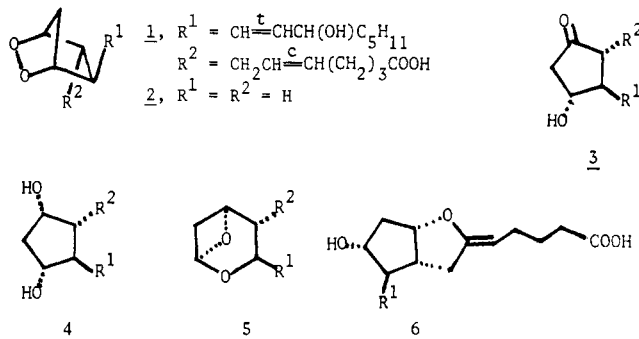
Department of Organic Chemistry, University of Groningen
Nijenborgh, 9747 AG Groningen, The Netherlands

Received December 27, 1978

Extraordinary Reactivity of the Prostaglandin Endoperoxide Nucleus. Nonpolar Rearrangement of 2,3-Dioxabicyclo[2.2.1]heptane and -[2.2.2]octane

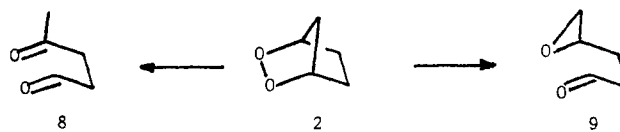
Sir:

Occasionally Nature provides us with molecules which not only have unusual structures, but which also exhibit extraordinary chemical reactivity. Prostaglandin (PG) endoperoxides¹ (e.g., **1**) possess an unusual bicyclic peroxide nucleus.^{2,2} They are a branch point in the oxidative transformation of polyunsaturated fatty acids into a vast array of physiologically active metabolites.³ The biological role of **1** depends in large measure on enzymatic conversion into prostaglandins (e.g., **3**, **4**), thromboxane A₂ (**5**),⁴ and prostacyclin (**6**).⁵ To provide a basis



for interpreting the complex biochemistry of **1**, we are studying the chemistry of the model endoperoxide **2** and homologues. We now report that the abnormally large solvent effects found for thermal decompositions of **2**⁶ are not observed for decomposition of the less strained homologue, 2,3-dioxabicyclo[2.2.2]octane (**7**).⁷ Furthermore, activation enthalpies and entropies for thermal decomposition of **2**, of the homologue **7**, and of *tert*-butyl peroxide in cyclohexane are remarkably different. ΔH^\ddagger increases with decreasing strain in the series.

Thermal decompositions of **2** and **7**⁷ were monitored by ¹H NMR. Relative rates in various solvents are listed in Table I. Both reactions follow first-order kinetics. As reported previously, the rate of decomposition of **2** increases with solvent polarity and is exceptionally rapid in protic solvents owing primarily to an extraordinary dependence of the rate of rearrangement to levulinaldehyde (**8**) on solvent polarity.⁶ The parallel first-order rearrangement of **2** to **9** is a nonpolar process which shows only a small dependence on solvent polarity.



In contrast, the rate of decomposition of **7** varies only slightly and erratically with changes in solvent polarity. The modest acceleration found for decomposition of **7** in protic solvents